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PTO/SB/08A (10-98)
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Substitute for form 1449A/PTO				Complete if Known	
INFORMATION DISCLOSURE STATEMENT BY APPLICANT (use as many sheets as necessary)				Application Number	Continuation of 10/028,547
				10/623431	
				Filing Date	July 18, 2003
				First Named Inventor	Jay D. Kranzler
				Group Art Unit	
Examiner Name					
Attorney Docket Number					CYPR 100 CIP CON
Sheet	1	of	4		

[illegible]

FOREIGN PATENT DOCUMENTS								
Examiner Initials*	Cite No. ¹	Foreign Patent Document			Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T ²
		Office. ³	Number ⁴	Kind Code ⁵ (if known)				
		EP	*0-759 299		Eli Lilly & Co.	02-26-1997		
		FR	*2-752 732		Pierre Fabre Medicament	03-06-1998		
N	WPCT	WO	96/22521		Asahi Kasei Kabushiki Kaisha	08-24-1995		
N	WPCT	WO	97/35574		Pierre Fabre Medicament	10-02-1997		
	WPCT	*WO	97/35584		Eli Lilly & Co.	10-02-1997		
N	WPCT	WO	98/08495		Pierre Fabre Medicament	03-05-1998		

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		Office. ³	Number ⁴	Kind Code ⁵ (if known)				
N		WO PCT	WO 98/36744		Pierre Fabre Medicament	08-27-1998		
		PCT	*WO 99/59593		Eli Lilly & Co.	11-25-1999		✓
		PCT	*WO 00/32178		Mueller	06-08-2000		✓
N		PCT	WO 01/26623		Laxdale Ltd	04-19-2001		✓
		PCT	*WO 02/053140		Pharmacia & Upjohn Co.	07-11-2002		✓

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		First Named Inventor	Jay D. Kranzler		
		Group Art Unit			
Examiner Name					
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OTHER ART -- NON PATENT LITERATURE DOCUMENTS			
Examiner's Initials*	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published	T ²
h		ARDID, et al., "Antidepressants and pain," La Lettre de Pharmacologie 13: 8 (1993).	
h		Cypress Bioscience, Inc., Investor Fact Sheet, August 2001.	
a		DRYSON, "Venlafaxine and fibromyalgia," NZ Med. J. 113(1105): 87 (2000).	
		*DWIGHT, et al., "An open clinical trial of venlafaxine treatment of fibromyalgia," Psychosomatics 39: 14-17 (1998). ✓	
		GOODNICK, et al., "Psychotropic treatment of chronic fatigue syndrome and related disorders," J. Clin. Psychiatry 13-20 (1993).	
		MEDLINE, et al., "Treatment of chronic fatigue syndrome with sibutramine," PCT Int'l Appl. 14 (09/28/2000).	
u		NAGAOKA, et al., "Beneficial effects of a serotonin-noradrenaline reuptake inhibitor on fibromyalgia syndrome: a case report," Med. Drug. J. 37: 10 (2001).	
		*NINAN, "Use of venlafaxine in other psychiatric disorders," Depression Anxiety 12S: 1:90-94 (2000). ✓	
n		NOGUCHI, et al., "Open channel block of NMDA receptors by conformationally restricted analogs of milnacipran and their protective effect against NMDA-induced neurotoxicity," Synapse 31: 87-96 (1999).	
		*NUTT & JOHNSON, "Potential applications of venlafaxine," Rev. Contemp. Pharmacother. 9: 321-331 (1998). ✓	

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Sheet	4	of	4
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OTHER ART -- NON PATENT LITERATURE DOCUMENTS			
Examiner's Initials*	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published	T ²
		*RAO, "The neuropharmacology of centrally-acting analgesic medications in fibromyalgia," Rheum. Dis. Clin. N. Amer. 28: 235-259 (2002).	
g		SHUTO, et al., "(±)-Z-2-(Aminomethyl)-1-phenylcyclopropanecarboximide derivatives as a new prototype of NMDA receptor antagonists," J. Med. Chem. 38: 2964-2968 (1995).	
n		SHUTO, et al., "(1S,2R)-1-(phenyl-2-((S)-1-aminopropyl))-N,N-diethylcyclopropanecarboxamide (PPDC), a new class of NMDA-receptor antagonist: molecular design by a novel conformational restriction strategy," Jpn. J. Pharmacol. 85: 207-213 (2001).	
u		SHUTO, et al., "Synthesis and biological activity of conformationally restricted analogs of milnacipran: (1S,2R)-1-(phenyl-2-((S)-1-aminopropyl))-N,N-diethylcyclopropanecarboxamide, an efficient noncompetitive N-methyl-D-aspartic acid receptor antagonist," J. Med. Chem. 39: 4844-4852 (1996).	
h		SHUTO, et al., "Synthesis and biological activity of conformationally restricted analogues of milnacipran: (1S,2R)-1-phenyl-2-((R)-1-amino-2-propynyl)-N,N-diethylcyclopropane-carboximide is a novel class of NMDA receptor channel blocker," J. Med. Chem. 41: 3507-3514 (1998).	

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